CLAIMS

What we claim is:

1. A compound of Formula (I):

$$Z \left\{ Y \right\}_{m}^{X} \left\{ W \right\}_{n}^{N}$$

$$(I)$$

wherein:

W and Y are independently a straight or branched chain C_{1-5} alkylene group optionally containing one double bond, one triple bond or carbonyl, wherein said C_{1-5} alkylene group is optionally substituted with halogen, hydroxyl, C_{1-4} alkyl, C_{1-4} haloalkyl or C_{1-4} alkoxy;

 $\label{eq:Xis-NR_3C(O)-, -C(O)NR_3, -NR_3S(O)_2-, -S(O)_2NR_3-, -NR_3C(O)NR_4-, -NR_3C(O)O-, -OC(O)NR_3-, -NR_3-, -C(O)-, -CH(OH)-, -C(NH)-, -O-, -S-, -S(O)- or -S(O)_2-;$

R₃ and R₄ are independently H, C₁₋₄ alkyl, phenyl or heteroaryl, wherein each of said alkyl, phenyl and heteroaryl are optionally substituted with 1 to 5 substituents selected from the group consisting of halogen, hydroxyl, thiol, cyano, nitro, C₁₋₄ haloalkyl, amino, C₁₋₄ alkylamino, di-C₁₋₄-alkylamino, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ haloalkoxy, C₁₋₄ alkylthio, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfonyl, C₁₋₄ haloalkylsulfinyl and C₁₋₄ haloalkylsulfonyl;

Z is H, halogen, phenyl or heteroaryl, wherein said phenyl and heteroaryl are optionally substituted with 1 to 5 substituents selected from the group consisting of halogen, hydroxy, thiol, cyano, nitro, C_{1-4} haloalkyl, amino, C_{1-4} alkylamino, di- C_{1-4} -alkylamino, C_{1-4} alkyl, C_{1-4} alkoxy, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} haloalkoxy, C_{1-4} alkylthio, C_{1-4} alkylsulfinyl, C_{1-4} haloalkylthio, C_{1-4} haloalkylsulfinyl and C_{1-4} haloalkylsulfonyl;

 R_1 is H, hydroxyl, halogen, $C_{1\!-\!4}$ alkyl or $C_{1\!-\!4}$ haloalkyl; R_2 is H or $C_{1\!-\!8}$ alkyl and

"n" and "m" are each independently 0 or 1; or a pharmaceutically acceptable salt, solvate or hydrate thereof; provided that:

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i)	when both R_1 and R_2 are H then $-[W]_n$ -X- $[Y]_m$ -Z together is not
CO ₂ H, C(O)-C	₆ H ₄ -p-O-C ₈ H ₁₇ , OCH ₂ CH ₃ , OH, CH ₂ CH ₂ CH ₂ CH ₂ CO ₂ H,
CH ₂ CH ₂ CH ₂ CO	D ₂ H, CH ₂ CO ₂ H and CH ₂ CH ₂ CO ₂ H:
•••	- • ····•

- ii) when R_1 is CH_3 and R_2 is H then $-[W]_n$ -X- $[Y]_m$ -Z together is not CH_2CO_2H , C(O)CH=CH C_6H_5 , $C(O)C_6H_4$ -p- OCH_3 , CO_2H , $C(O)CH_3$, $C(O)C_6H_4$ -o- CH_3 , $C(O)C_6H_4$ -o-CI, and $C(O)C_6H_5$;
- iii) when R_1 is Br and R_2 is H then $-[W]_n$ -X- $[Y]_m$ -Z together is not CO_2H ;
- iv) when R_1 is OH and R_2 is H then $-[W]_n$ -X- $[Y]_m$ -Z together is not CO_2H ;
- v) when R₁ is H and R₂ is CH₃ then –[W]_n-X-[Y]_m-Z together is not 2,6-dichloro-4-trifluoromethylphenoxy, C(O)NH-C₆H₄-p-OCH₂CH₃, NHC(O)CH(CH₃)₂, SCH₃, C(O)-C₆H₄-p-O-C₈H₁₇, SCH₂CH₃, C(O)NHC₆H₅, CH(OCH₃)₂, CH₂OC(O)CH₃, CO₂H, CO₂CH₃, C(O)C₆H₄-p-NO₂, C(O)C₆H₅, CH₂CH₂CO₂CH₃, CH₂CH₂CO₂CH₃, CH₂CH₂CO₂CH₃, CH₂CH₂CO₂CH₃, CH₂CH₂CO₂CH₃, and CH₂CO₂CH₃;
- vi) when R_1 is OH and R_2 is CH₃ then $-[W]_n$ -X- $[Y]_m$ -Z together is not CH₂OCH₂C₆H₅, CH₂OCH(CH₃)₂ and CH₂OH;
 - vii) when R₂ is CH₃ then:

 R_1 is not CH_3 and $-[W]_n$ -X- $[Y]_m$ -Z together is not 2,6-dichloro-4-trifluoromethylphenoxy;

 R_1 is not I and $-[W]_n$ -X- $[Y]_m$ -Z together is not $CO_2C(CH_3)_3$; R_1 is not $C(CH_3)_3$ and $-[W]_n$ -X- $[Y]_m$ -Z together is not formyl;

 R_1 is not Br and $-[W]_n$ -X- $[Y]_m$ -Z together is not CO_2CH_3 ; and

 R_1 is not $CH_2CH_2CH_2CH_3$ and $-[W]_n\text{-}X\text{-}[Y]_m\text{-}Z$ together is not formyl;

- viii) when R₁ is H and R₂ is CH₂CH₃ then -[W]_n-X-[Y]_m-Z together is not CH₂SCH₂CH₃, OCH₂CH₂CH=CH₂, CH₂CH₂CH₂OH, CH₂CH₂CHO, CO₂CH₂CH₃, OCH₃, C(O)CH₂Br, CO₂C₈H₁7, formyl, OH, CH₂N(CH₂CH₂Cl)₂, CH(CH₃)OC(O)CH₃, CH₂OH, CH₂OC(O)CH₃, C(O)CH₃, C(O)C₆H₅ and C(O)NHCH₂CO₂CH₂CH₃.
- ix) when R_1 is CH_3 and R_2 is CH_2CH_3 then $-[W]_n-X-[Y]_m-Z$ together is not $CH(OH)C_6H_4-p-N(CH_3)_2$, $C(O)CH_2C(O)CH_3$, $CO_2CH_2C_6H_5$, CO_2CH_3 , $C(O)CH_2CH_2CH_3$, $C(O)CH_3$, $C(O)C_6H_4-p-OCH_3$, $C(O)C_6H_4-o-Br$, $C(O)C_6H_4-o-Cl$, $C(O)C_6H_4-o-Cl$, $C(O)C_6H_5$ and $C(O)C_6H_5$;
 - x) when R₂ is CH₂CH₃ then:

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 R_1 is not I and $-[W]_n$ -X- $[Y]_m$ -Z together is not $CO_2CH_2CH_3$; R_1 is not CF_3 and $-[W]_n$ -X- $[Y]_m$ -Z together is not $CO_2CH_2CH_3$; and

R₁ is not Br and -[W]_n-X-[Y]_m-Z together is not CO₂CH₂CH₃;

- xi) when R_1 is OH and R_2 is CH_2CH_3 then $-[W]_n$ -X- $[Y]_m$ -Z together is not $C(O)C_6H_5$, $C(O)NH_2$ and $CO_2CH_2CH_3$;
- xii) when R_1 is H and R_2 is $C(CH_3)_3$ then $-[W]_n$ -X- $[Y]_m$ -Z together is not $CO_2C(CH_3)_3$, $C(O)NHC(O)CH_3$ and $C(O)NH_2$;
- xiii) when R_1 is OH and R_2 is $CH_2CH_2CH_2CH_3$ then $-[W]_n$ -X- $[Y]_m$ -Z together is not $C(O)C_6H_5$; and
 - xiv) when X is -NR₃- then "n" is 1.
- 2. The compound according to claim 1 wherein "n" is 0.
- 3. The compound according to claim 1 wherein "n" is 1.
- 4. The compound according to any one of claims 1 to 3 wherein "m" is 0.
- 20 5. The compound according to any one of claims 1 to 3 wherein "m" is 1.
 - 6. The compound according to any one of claims 1, 3, 4 and 5 wherein W is the straight or branched C₁₋₅ alkylene group optionally containing one double bond, one triple bond or carbonyl, wherein said C₁₋₅ alkylene group is optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 7. The compound according to claim 6 wherein W is -CH₂- optionally substituted with halogen, hydroxyl, C_{1-4} alkyl or C_{1-4} alkoxy.
- 30 8. The compound according to claim 7 wherein W is -CH(CH₃)- optionally substituted with halogen, hydroxyl or C₁₋₄ alkoxy.
 - 9. The compound according to claim 7 wherein W is -C(CH₃)₂-.
- The compound according to claim 6 wherein W is -CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.

- 11. The compound according to claim 10 wherein W is -CH(CH₃)CH₂- or -CH₂CH(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- The compound according to claim 10 wherein W is -C(CH₃)₂CH₂- or -CH₂C(CH₃)₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- The compound according to claim 10 wherein W is -CH(OCH₃)CH₂- or
 -CH₂CH(OCH₃)- optionally substituted with halogen, hydroxyl or C₁₋₄ alkyl.
 - 14. The compound according to claim 6 wherein W is -CH₂CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 15. The compound according to claim 6 wherein W is -CH₂CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 16. The compound according to claim 6 wherein W is -CH=CH- optionally substituted with C_{1-4} alkyl or C_{1-4} alkoxy.
 - 17. The compound according to claim 6 wherein W is -C = C.

- 18. The compound according to claim 6 wherein W is -C(O)-.
- The compound according to claim 6 wherein W is -CH₂C(O)- or -C(O)CH₂- optionally substituted with halogen, hydroxyl, C_{1-4} alkyl or C_{1-4} alkoxy.
 - 20. The compound according to claim 19 wherein W is -CH(CH₃)C(O)- or -C(O)CH(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - The compound according to claim 19 wherein W is -C(CH₃)₂C(O)- or -C(O)C(CH₃)₂-.
- The compound according to claim 6 wherein W is -CH₂CH₂C(O)- or -C(O)CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.

- 23. The compound according to claim 22 wherein W is -C(CH₃)₂CH₂C(O)- or -C(O)CH₂C(CH₃)₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 24. The compound according to claim 6 wherein W is -CH₂C(O)CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.

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- The compound according to claim 6 wherein W is -CH₂CH₂CH₂C(O)- or
 -C(O)CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 26. The compound according to claim 6 wherein W is -CH(CH₃)CH₂CH₂C(O)- or -C(O)CH₂CH₂CH(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 27. The compound according to claim 6 wherein W is -CH₂CH₂C(O)CH₂- or -CH₂C(O)CH₂-CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 28. The compound according to claim 6 wherein W is -CH=CHC(O)- or -C(O)CH=CH- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 25 29. The compound according to claim 6 wherein W is -C(CH₃)=CHC(O)- or -C(O)CH=C(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 30. The compound according to any one of claims 1, 2, 3 and 5 to 28 wherein Y is the straight or branched chain C₁₋₅ alkylene group optionally containing one double bond, one triple bond or carbonyl, wherein said C₁₋₅ alkylene group is optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- The compound according to claim 30 wherein Y is -CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 32. The compound according to claim 31 wherein Y is -CH(CH₃)- optionally substituted

with halogen, hydroxyl or C_{1-4} alkoxy.

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- 33. The compound according to claim 31 wherein Y is -C(CH₃)₂-.
- 5 34. The compound according to claim 30 wherein Y is -CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - The compound according to claim 34 wherein Y is -CH(CH₃)CH₂- or -CH₂CH(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 36. The compound according to claim 34 wherein Y is -C(CH₃)₂CH₂- or -CH₂C(CH₃)₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- The compound according to claim 34 wherein Y is -CH(OCH₃)CH₂- or
 -CH₂CH(OCH₃)- optionally substituted with halogen, hydroxyl or C₁₋₄ alkyl.
- The compound according to claim 30 wherein Y is -CH₂CH₂-Optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 39. The compound according to claim 30 wherein Y is -CH₂CH₂CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 25 40. The compound according to claim 30 wherein Y is -CH=CH- optionally substituted with C_{1-4} alkyl or C_{1-4} alkoxy.
 - 41. The compound according to claim 30 wherein Y is -C ≡C-.
- 30 42. The compound according to claim 30 wherein Y is $-C = CCH_2$ or $-CH_2C = C$ optionally substituted with halogen, hydroxyl, C_{1-4} alkyl or C_{1-4} alkoxy.
 - 43. The compound according to claim 30 wherein Y is -C(O)-.
- The compound according to claim 30 wherein Y is -CH₂C(O)- or -C(O)CH₂-optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.

- 45. The compound according to claim 44 wherein Y is -CH(CH₃)C(O)- or -C(O)CH(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 5 46. The compound according to claim 44 wherein Y is -C(CH₃)₂C(O)- or -C(O)C(CH₃)₂-.

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- 47. The compound according to claim 30 wherein Y is -CH₂CH₂C(O)- or -C(O)CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 48. The compound according to claim 47 wherein Y is -C(CH₃)₂CH₂C(O)- or -C(O)CH₂C(CH₃)₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- 49. The compound according to claim 30 wherein Y is -CH₂C(O)CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- The compound according to claim 30 wherein Y is -CH₂CH₂CH₂C(O)- or -C(O)CH₂CH₂CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 51. The compound according to claim 30 wherein Y is -CH(CH₃)CH₂CH₂C(O)- or -C(O)CH₂CH₂CH(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 52. The compound according to claim 30 wherein Y is -CH₂CH₂C(O)CH₂- or -CH₂C(O)CH₂- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
 - 53. The compound according to claim 30 wherein Y is -CH=CHC(O)- or -C(O)CH=CH- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.
- The compound according to claim 30 wherein Y is -C(CH₃)=CHC(O)- or -C(O)CH=C(CH₃)- optionally substituted with halogen, hydroxyl, C₁₋₄ alkyl or C₁₋₄ alkoxy.

- 55. The compound according to any one of claims 1 to 54 wherein X is -NR₃C(O)-.
- 56. The compound according to any one of claims 1 to 54 wherein X is -C(O)NR₃-.

57. The compound according to any one of claims 1 to 54 wherein X is -NR₃S(O)₂-.

58. The compound according to any one of claims 1 to 54 wherein X is -S(O)₂NR₃-.

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- 59. The compound according to any one of claims 1 to 54 wherein X is -NR₃C(O)NR₄-.
- 15 60. The compound according to any one of claims 1 to 54 wherein X is -NR₃C(O)O-.
 - 61. The compound according to any one of claims 1 to 54 wherein X is -OC(O)NR₃-.
 - 62. The compound according to any one of claims 1 to 54 wherein X is -NR₃-.
 - 63. The compound according to any one of claims 55 to 62 wherein R₃ is H or CH₃.
 - 64. The compound according to claim 59 wherein R₄ is H or CH₃.
- 25 65. The compound according to any one of claims 1 to 54 wherein X is -C(O)-.
 - 66. The compound according to any one of claims 1 to 54 wherein X is -CH(OH)-.
- 67. The compound according to any one of claims 1 to 54 wherein X is -C(NH)-.
 - 68. The compound according to any one of claims 1 to 54 wherein X is -O-.
 - 69. The compound according to any one of claims 1 to 54 wherein X is -S-.
- The compound according to any one of claims 1 to 54 wherein X is -S(O)-.
 - 71. The compound according to any one of claims 1 to 54 wherein X is $-S(O)_2$.

72. The compound according to any one of claims 1 to 71 wherein Z is H.

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- 73. The compound according to any one of claims 1 to 71 wherein Z is halogen.
- 74. The compound according to any one of claims 1 to 71 wherein Z is phenyl.
- 75. The compound according to claim 74 wherein the phenyl is optionally substituted with 1 to 5 substituents selected from the group consisting of halogen, C₁₋₄ haloalkyl, C₁₋₄ alkylamino, di-C₁₋₄-alkylamino, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfonyl, C₁₋₄ haloalkylthio, C₁₋₄ haloalkylsulfinyl and C₁₋₄ haloalkylsulfonyl.
- 76. The compound according to claim 75 wherein the phenyl is optionally substituted with 1 to 3 substituents selected from the group consisting of -F, -Cl, -Br, -CF₃, -NHCH₃, -N(CH₃)₂, -CH₃, -CH₂CH₃, -OCH₃ and -OCF₃.
 - 77. The compound according to any one of claims 1 to 71 wherein Z is heteroaryl.
- 78. The compound according to claim 77 wherein the heteroaryl is optionally substituted with 1 to 5 substituents selected from the group consisting of halogen, C₁₋₄ haloalkyl, C₁₋₄ alkylamino, di-C₁₋₄-alkylamino, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, C₁₋₄ alkylthio, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfonyl, C₁₋₄ haloalkylthio, C₁₋₄ haloalkylsulfinyl and C₁₋₄ haloalkylsulfonyl.
 - 79. The compound according to claim 78 wherein the phenyl is optionally substituted with 1 to 3 substituents selected from the group consisting of -F, -Cl, -Br, -CF₃, -NHCH₃, -N(CH₃)₂, -CH₃, -CH₂CH₃, -OCH₃ and -OCF₃.
- 30 80. The compound according to any one of claims 1 to 79 wherein R_1 is H.
 - 81. The compound according to any one of claims 1 to 79 wherein R_1 is hydroxyl.
 - 82. The compound according to any one of claims 1 to 78 wherein R_1 is halogen.
 - 83. The compound according to any one of claims 1 to 78 wherein R_1 is C_{1-4} alkyl.

- 84. The compound according to any one of claims 1 to 78 wherein R₁ is C₁₋₄ haloalkyl.
- 85. The compound according to any one of claims 1 to 84 wherein R₂ is H.
- 5 86. The compound according to any one of claims 1 to 84 wherein R₂ is C₁₋₈ alkyl.
 - 87. A pharmaceutical composition comprising a pharmaceutically acceptable carrier in combination with at least one compound according to Formula (I):

10 wherein:

W and Y are independently a straight or branched chain C_{1-5} alkylene group optionally containing one double bond, one triple bond or carbonyl, wherein said C_{1-5} alkylene group is optionally substituted with halogen, hydroxyl, C_{1-4} alkyl, C_{1-4} haloalkyl or C_{1-4} alkoxy;

 $\label{eq:Xis-NR3C(O)-,-C(O)NR3,-NR3S(O)2-,-S(O)2NR3-,-NR3C(O)NR4-,-NR3C(O)O-,-OC(O)NR3-,-NR3-,-C(O)-,-CH(OH)-,-C(NH)-,-O-,-S-,-S(O)- or -S(O)2-;}$

R₃ and R₄ are independently H, C₁₋₄ alkyl, phenyl or heteroaryl, wherein each of said alkyl, phenyl and heteroaryl are optionally substituted with 1 to 5 substituents selected from the group consisting of halogen, hydroxyl, thiol, cyano, nitro, C₁₋₄ haloalkyl, amino, C₁₋₄ alkylamino, di-C₁₋₄-alkylamino, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ haloalkoxy, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfonyl, C₁₋₄ haloalkylsulfinyl and C₁₋₄ haloalkylsulfonyl;

Z is H, halogen, phenyl or heteroaryl, wherein said phenyl and heteroaryl are optionally substituted with 1 to 5 substituents selected from the group consisting of halogen, hydroxy, thiol, cyano, nitro, C₁₋₄ haloalkyl, amino, C₁₋₄ alkylamino, di-C₁₋₄-alkylamino, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ haloalkoxy, C₁₋₄ alkylthio, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfinyl, C₁₋₄ haloalkylsulfinyl and C₁₋₄ haloalkylsulfonyl;

R₁ is H, hydroxyl, halogen, C₁₋₄ alkyl or C₁₋₄ haloalkyl;

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R₂ is H or C₁₋₈ alkyl and "n" and "m" are each independently 0 or 1; or a pharmaceutically acceptable salt, solvate or hydrate thereof; provided that when X is -NR3- then "n" is 1.

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88. The pharmaceutical composition according to claim 87 further comprising one or more agents selected from the group consisting of α -glucosidase inhibitor, aldose reductase inhibitor, biguanide, HMG-CoA reductase inhibitor, squalene synthesis inhibitor, fibrate, LDL catabolism enhancer, angiotensin converting enzyme inhibitor, insulin secretion enhancer and thiazolidinedione.

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89. The pharmaceutical composition according to claim 88 wherein the agent is a α glucosidase inhibitor.

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90. The pharmaceutical composition according to claim 89 wherein the α -glucosidase inhibitor is acarbose, voglibose or miglitol.

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The pharmaceutical composition according to claim 90 wherein the α -glucosidase 91. inhibitor is voglibose.

The pharmaceutical composition according to claim 88 wherein the agent is an aldose 92. reductase inhibitor.

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93. The pharmaceutical composition according to claim 92 wherein the aldose reductase inhibitor is tolurestat; epalrestat; imirestat; zenarestat; zopolrestat; or sorbinil.

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The pharmaceutical composition according to claim 88 wherein the agent is a biguanide.

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The pharmaceutical composition according to claim 94 wherein the biguanide is 95. phenformin, metformin or buformin.

96. The pharmaceutical composition according to claim 95 wherein the biguanide is metformin.

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97. The pharmaceutical composition according to claim 88 wherein the agent is a HMG- CoA reductase inhibitor.

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- 98. The pharmaceutical composition according to claim 97 wherein the HMG-CoA reductase inhibitor is rosuvastatin, pravastatin, simvastatin, lovastatin, atorvastatin, fluvastatin or cerivastatin.
- 99. The pharmaceutical composition according to claim 88 wherein the agent is a fibrate.
- 100. The pharmaceutical composition according to claim 99 wherein the fibrate is
 bezafibrate, beclobrate, binifibrate, ciplofibrate, clinofibrate, clofibrate, clofibric acid,
 etofibrate, fenofibrate, gemfibrozil, nicofibrate, pirifibrate, ronifibrate, simfibrate, or
 theofibrate.
- The pharmaceutical composition according to claim 88 wherein the agent is an angiotensin converting enzyme inhibitor.
 - 102. The pharmaceutical composition according to claim 101 wherein the angiotensin converting enzyme inhibitor is captopril, enalapril, alacepril, delapril; ramipril, lisinopril, imidapril, benazepril, ceronapril, cilazapril, enalaprilat, fosinopril, moveltopril, perindopril, quinapril, spirapril, temocapril or trandolapril.
 - 103. The pharmaceutical composition according to claim 88 wherein the agent is an insulin secretion enhancer.
- The pharmaceutical composition according to claim 103 wherein the insulin secretion enhancer is tolbutamide; chlorpropamide; tolazamide; acetohexamide; glycopyramide; glibenclamide; gliclazide; 1-butyl-3-metanilylurea; carbutamide; glibonuride; glipizide; gliquidone; glisoxepid; glybuthiazole; glibuzole; glyhexamide; glymidine; glypinamide; phenbutamide; tolcyclamide, glimepiride, nateglinide, or mitiglinide.
 - 105. The pharmaceutical composition according to claim 88 wherein the agent is a thiazolidinedione.
- The pharmaceutical composition according to claim 105 wherein the thiazolidinedione is rosiglitazone or pioglitazone.

107. The pharmaceutical composition according to claim 106 wherein the thiazolidinedione is rosiglitazone.

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- 108. The compound according to any one of claims 1 to 86 for use in a method of treatment of the human or animal body by therapy.
- 109. The compound according to any one of claims 1 to 86 for use in a method of prophylaxis or treatment of a metabolic-related disorder of the human or animal body by therapy.
- 110. A method for prophylaxis or treatment of a metabolic-related disorder in an individual in need of said prophylaxis or treatment comprising administering to the individual a therapeutically effective amount of a compound according to any one of claims 1 to 86 or a pharmaceutical composition according to any one of claims 87 to 107.
- 111. A method of modulating a RUP25 receptor in an individual comprising contacting the receptor with a compound according to any one of claims 1 to 86.
- 20 112. The method of modulating the RUP25 receptor according to claim 111 wherein the compound is an agonist.
 - 113. The method of modulating the RUP25 receptor according to claim 111 or 112 wherein the modulation of the RUP25 receptor is for prophylaxis or treatment of a metabolic-related disorder in an individual in need of said prophylaxis or treatment.
 - 114. The method according to claim 110 or 113 wherein the metabolic-related disorder is selected from the group consisting of dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance, obesity, impaired glucose tolerance, atheromatous disease, hypertension, stroke, Syndrome X, heart disease and type 2 diabetes.
 - 115. The method according to claim 114 wherein the metabolic-related disorder is dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance and type 2 diabetes.
 - 116. The method according to claim 115 wherein the metabolic-related disorder is dyslipidemia.

- 117. The method according to claim 115 wherein the metabolic-related disorder is atherosclerosis.
- 5 118. The method according to claim 115 wherein the metabolic-related disorder is coronary heart disease.

- 119. The method according to claim 115 wherein the metabolic-related disorder is insulin resistance.
- 120. The method according to claim 115 wherein the metabolic-related disorder is type 2 diabetes.
- Use of a compound according to any one of claims 1 to 86 for production of a medicament for use in prophylaxis or treatment of a metabolic-related disorder.
 - 122. The use according to claim 121 further comprising one or more agents selected from the group consisting of α-glucosidase inhibitor, aldose reductase inhibitor, biguanide, HMG-CoA reductase inhibitor, squalene synthesis inhibitor, fibrate, LDL catabolism enhancer, angiotensin converting enzyme inhibitor, insulin secretion enhancer and thiazolidinedione.
 - 123. The use according to claim 122 wherein the agent is a α -glucosidase inhibitor.
- 25 124. The use according to claim 123 wherein the α -glucosidase inhibitor is acarbose, voglibose or miglitol.
 - 125. The use according to claim 124 wherein the α -glucosidase inhibitor is voglibose.
- The use according to claim 122 wherein the agent is an aldose reductase inhibitor.
 - 127. The use according to claim 126 wherein the aldose reductase inhibitor is tolurestat; epalrestat; imirestat; zenarestat; zopolrestat; or sorbinil.
- The use according to claim 122 wherein the agent is a biguanide.

- 129. The use according to claim 128 wherein the biguanide is phenformin, metformin or buformin.
- 130. The use according to claim 129 wherein the biguanide is metformin.

The use according to claim 122 wherein the agent is a HMG-CoA reductase inhibitor.

- 132. The use according to claim 131 wherein the HMG-CoA reductase inhibitor is rosuvastatin, pravastatin, simvastatin, lovastatin, atorvastatin, fluvastatin or cerivastatin.
- 133. The use according to claim 122 wherein the agent is a fibrate.

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- The use according to claim 133 wherein the fibrate is bezafibrate, beclobrate, binifibrate, ciplofibrate, clinofibrate, clofibrate, clofibrate acid, etofibrate, fenofibrate, gemfibrozil, nicofibrate, pirifibrate, ronifibrate, simfibrate, or theofibrate.
 - 135. The use according to claim 122 wherein the agent is an angiotensin converting enzyme inhibitor.
 - 136. The use according to claim 135 wherein the angiotensin converting enzyme inhibitor is captopril, enalapril, alacepril, delapril; ramipril, lisinopril, imidapril, benazepril, ceronapril, cilazapril, enalaprilat, fosinopril, moveltopril, perindopril, quinapril, spirapril, temocapril or trandolapril.
 - 137. The use according to claim 122 wherein the agent is an insulin secretion enhancer.
 - 138. The use according to claim 137 wherein the insulin secretion enhancer is tolbutamide; chlorpropamide; tolazamide; acetohexamide; glycopyramide; glibenclamide; gliclazide; 1-butyl-3-metanilylurea; carbutamide; glibonuride; glipizide; gliquidone; glisoxepid; glybuthiazole; glibuzole; glyhexamide; glymidine; glypinamide; phenbutamide; tolcyclamide, glimepiride, nateglinide, or mitiglinide.
 - 139. The use according to claim 122 wherein the agent is a thiazolidinedione.
 - 140. The use according to claim 139 wherein the thiazolidinedione is rosiglitazone or pioglitazone.

- 141. The use according to claim 140 wherein the thiazolidinedione is rosiglitazone.
- The use according to any one of claims 121 to 141 wherein the metabolic-related disorder is dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance, obesity, impaired glucose tolerance, atheromatous disease, hypertension, stroke, Syndrome X, heart disease and type 2 diabetes.
- The use according to claim 142 wherein the metabolic-related disorder is dyslipidemia, atherosclerosis, coronary heart disease, insulin resistance and type 2 diabetes.
 - 144. The use according to claim 143 wherein the metabolic-related disorder is dyslipidemia.

- 145. The use according to claim 143 wherein the metabolic-related disorder is atherosclerosis.
- The use according to claim 143 wherein the metabolic-related disorder is coronary heart disease.
 - 147. The use according to claim 143 wherein the metabolic-related disorder is insulin resistance.
- 25 The use according to claim 143 wherein the metabolic-related disorder is type 2 diabetes.
- The method of producing a pharmaceutical composition comprising admixing at least one compound according to any one of claims 1 to 86 and a pharmaceutically acceptable carrier or excipient.